Date of Report	23/10/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. HARSHITA	Patient ID	012210220099
Birthday	15/08/1993	Sample ID	11348013
Age at term	29.08	Sample Date	22/10/2022
Gestational age	11+3		

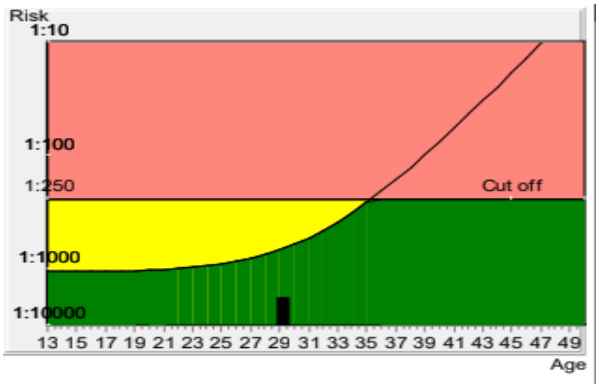
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	3.1 mIU/ml	0.87
fb-hCG	21.5 ng/ml	0.44

Risks at sampling date	
Age Risk	1:680
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	11+2
Method	CRL (<>Robinson)
Scan date	21/10/2022
Nuchal translucency	1.1
Nuchal translucency MoM	0.88
Nasal bone	Present

Risk



Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21with NT test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off Risk above Age Risk Risk below Age risk